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(54) **Poly(aryl ether sulfone) compositions.**

(57) A poly(aryl ether sulfone) blend comprising (a) from about 25 to about 99 percent by weight of a poly(biphenyl ether sulfone) and (b) from about 1 to about 75 percent by weight of a second poly(aryl ether sulfone) comprising bisphenol A moieties.

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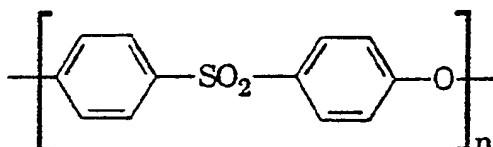
This invention is directed to immiscible poly(aryl ether sulfone) blends possessing improved properties comprising (a) from about 25 to about 99 percent by weight of a poly(biphenyl ether sulfone) preferably 60 to 99 weight percent and (b) from about 1 to about 75 percent by weight of a second poly(aryl ether sulfone) comprising bisphenol A moieties preferably 40 to 1 weight percent. The compositions described herein, particularly the preferred compositions, retain the very attractive characteristics of their constituents, and display excellent mechanical properties and heat resistance.

Articles made from these poly(aryl ether sulfones) containing at least about 35 to 99 weight percent poly(biphenyl ether sulfone) can be steam-sterilized while under stresses of 500 psi or greater, moreover they are not affected by corrosion-reducing additives such as morpholine, for example. Also, the above materials demonstrate good chemical resistance in contact with commonly used hospital cleaners and detergents.

Poly(aryl ether sulfones) have been known for about two decades. They are tough linear polymers that possess a number of attractive features such as excellent high temperature resistance, good electrical properties, and very good hydrolytic stability. At least three poly(aryl ether sulfones) are commercially available. A poly(aryl ether sulfone) is available from Imperial Chemical Industries, Ltd. It has the formula (1)

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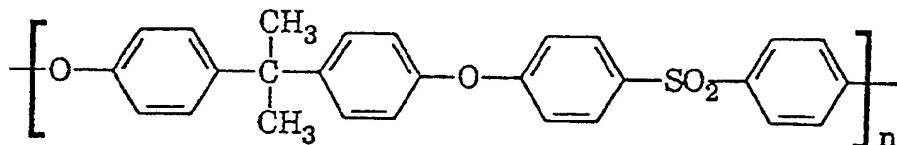
(1)

and is produced by the polycondensation of 4,4'-dihydroxydiphenyl sulfone with 4,4'-dichlorodiphenyl sulfone as described in, for example, Canadian Patent No. 847,963. The polymer contains no aliphatic moieties and has a Tg of approximately 220°C.

Another commercial poly(aryl ether sulfone) is available from Amoco Performance Products, Inc., under the trademark of UDEL®. It corresponds to formula (2),

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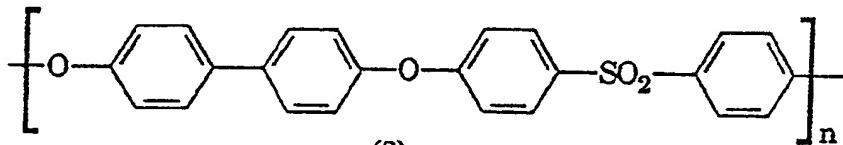
(2)

has a Tg of about 190°C, and is made via the nucleophilic polycondensation of bisphenol-A di-sodium salt with 4,4'-dichlorodiphenyl sulfone, as described in U.S. Patent No. 4,108,837.

A third commercial poly(aryl ether sulfone) is also available from Amoco Performance Products, Inc., under the trademark of Radel® R. It corresponds to formula (3) has a Tg of about 220°C, and is produced by the polycondensation of biphenol with 4,4'-dichlorodiphenyl sulfone as described in, for example, Canadian Patent No. 847,963.

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(3)

Over the years, there has developed a substantial body of patent and other literature directed to the formation and properties of poly(aryl ether sulfones and other poly(aryl ethers) (all hereinafter called "PAE"). A broad range of PAE's was achieved by Johnson et al., J. of Polymer Science, A-1, Vol. 5, 1967, pp. 2415-2427; Johnson et al., U.S. Patent Nos. 4,108,837 and 4,175,175. Johnson et al. show that a very broad range of PAE's can be formed by the nucleophilic aromatic substitution (condensation) reaction of an activated aromatic dihalide and an aromatic diol. By this method, Johnson et al. created a host of new PAE's.

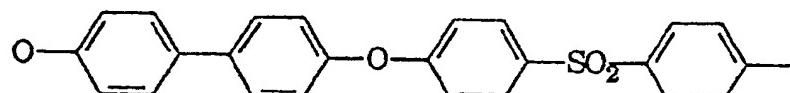
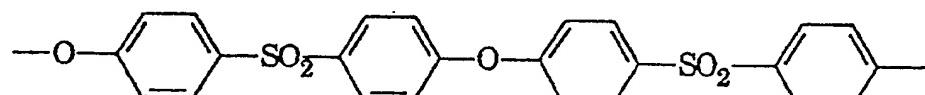
Because of their excellent mechanical and thermal properties, coupled with outstanding hydrolytic stability,

the poly(aryl ether sulfones) have been utilized in the medical market for a variety of purposes for at least ten years. These medical devices constitute a wide variety of articles. Obviously, one of the major attributes of the poly(aryl ether sulfones) is their ability to be steam autoclaved repeatedly without loss of properties. Steam autoclaving is a very severe test, requiring both high temperature and hydrolytic stability, and involving cyclical effects - wet/dry, hot/cold.

The poly(aryl ether sulfones) (1) and (2) show some important deficiencies, however. Indeed, parts molded from these materials, stress-crack when steam sterilized under stresses of say 500 psi or greater when excessive concentration of boiler additives, such as morpholine are employed to reduce corrosion in the steam generating system; or, when in contact with commonly used hospital cleaners and detergents.

While poly(biphenyl ether sulfone) (3) and parts molded therefrom have substantially better properties than poly(aryl ether sulfones) (1) and (2) it is substantially more expensive than (1) and (2) due to the high cost of biphenol.

British Patent Application No. 2,088,396 describes copolymers containing units (4) and (5):



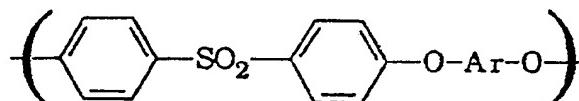
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(5)

The claimed copolymers comprise about 80 to 10 mole percent of repeat units (4), and correspondingly about 20 to 90 mole percent of repeat units (5). The application states that the incorporation of (5) into the poly(aryl ether sulfone) (1) yields materials with improved resistance to hot water crazing. The application does not mention steam-sterilizability under load; nor does it teach that the copolymers show resistance to stress-cracking in the presence of boiler additives such as morpholine.

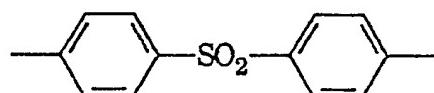
The general object of this invention is to provide blends of poly(biphenyl ether sulfones) having many of the base properties of the poly(biphenyl ether sulfone). Another object of this invention is to provide medical articles from blends of poly(biphenyl ether sulfones) which can be steam-sterilized while under stresses of 500 psi or greater without stresscracking even in the presence of morpholine. Other objects appear hereafter.

The general objects of this invention can be attained with immiscible blends comprising (a) from about 25 to about 99 percent by weight of a poly(biphenyl ether sulfone) and (b) from about 1 to 75 percent by weight of a second poly(aryl ether sulfone) comprising bisphenol A. Other things being equal blends of polyarylether (2) and poly(aryl ether sulfone) (3) wherein poly(aryl ether sulfone) (3) comprises at least 60% of the two polymers have substantially the same properties as the more expensive poly(aryl ether sulfone) (3). The two polymers can also be used in weight percent ratios of poly(biphenyl ether sulfone) of from about 50 to 99 to poly(aryl ether sulfone) (2) of about 50 to 1 weight percent.

Briefly, the poly(biphenyl ether sulfones) useful in this invention comprise the repeating unit



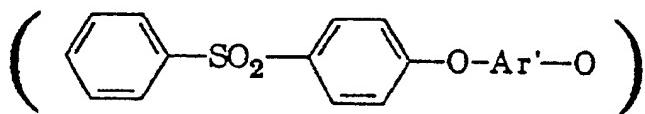
50 whereon at least 50 and preferably at least 75 mole percent of the divalent Ar groups are p-biphenylene groups and the remainder (0 to 50 mole percent) at least one member selected from the group consisting of p-phenylene,



, etc. In general, the higher the concentration of biphenyl or biphenylene groups the better the properties of the polymer.

The poly(aryl ether sulfones) comprising Bisphenol A moieties useful in this invention comprise the repeating unit

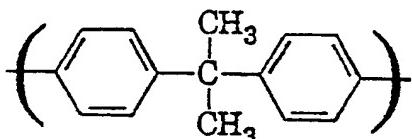
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wherein at least 50 and preferably at least 75 mole percent of the divalent Ar' groups are bisphenol A moieties

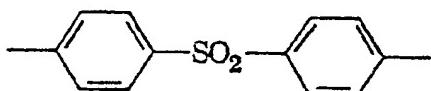
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and the remainder (0 to 50 mole percent) at least one member selected from the group consisting of p-phenylene and

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The poly(aryl ether sulfones) can be prepared by either of two methods, i.e., the carbonate method or the alkali metal hydroxide method.

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In the carbonate method, the polymers are prepared by contacting substantially equimolar amounts of the hydroxy-containing compounds and dihalodiarylsulfones, e.g., 4,4'-dichlorodiphenyl sulfone or 4,4'-difluorodiphenyl sulfone, with from about 0.5 to about 1.0 mole of an alkali metal carbonate per mole of hydroxyl group in a solvent mixture comprising a solvent which forms an azeotrope with water in order to maintain the reaction medium at substantially anhydrous conditions during the polymerization.

The temperature of the reaction mixture is kept at about 170°C to about 250°C, preferably from about 210°C to about 235°C for about one to 15 hours.

40

In a modification which is particularly suitable for making copolymers from bisphenol A and one or more additional dihydroxy compounds, the reactants other than said additional dihydroxy compounds are charged and heated at from about 120°C to about 180°C for about one to about 5 hours, said additional dihydroxy compounds are added, the temperature is raised and the mixture is heated at from about 200°C to about 250°C, preferably from about 210°C to about 240°C, for about one to 10 hours. This modification is further described in the copending U.S. Patent application of Donald R. Kelsey, et al., Serial No. 068,973, filed July 1, 1987, commonly assigned.

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The reaction is carried out in an inert atmosphere, e.g., nitrogen, at atmospheric pressure, although higher or lower pressures may also be used.

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The polyarylethersulfone is then recovered by conventional techniques such as coagulation, solvent evaporation, and the like. The solvent mixture comprises a solvent which forms an azeotrope with water and a polar aprotic solvent. The solvent which forms an azeotrope with water includes an aromatic hydrocarbon such as benzene, toluene, xylene, ethylbenzene, chlorobenzene, and the like.

The polar aprotic solvents employed in this invention are those generally known in the art for the manufacture of polyarylether sulfones and include sulfur containing solvents such as those of the formula:

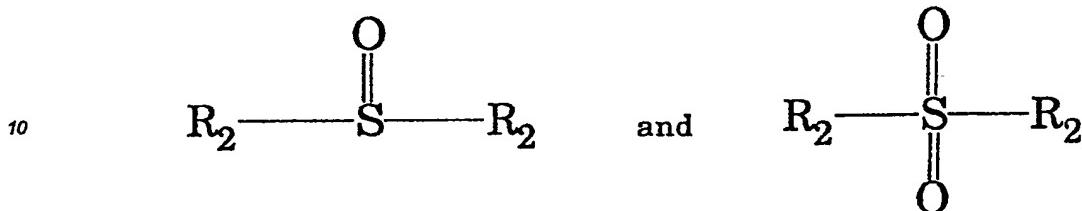
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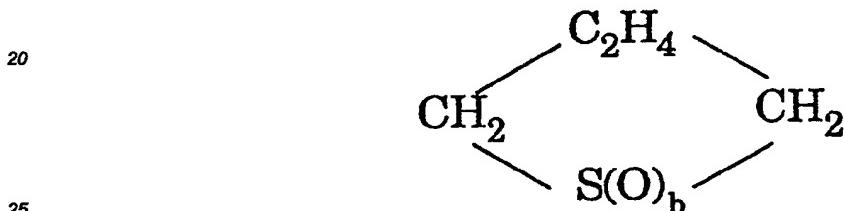
in which each R₁ represents a monovalent lower hydrocarbon group free of aliphatic unsaturation, which pref-

erably contains less than about 8 carbon atoms or when connected together represents a divalent alkylene group with b being an integer from 1 to 2 inclusive. Thus, in all of these solvents, all oxygens and two carbon atoms are bonded to the sulfur atom. Contemplated for use in this invention are such solvents as those having the formula:

5



15 where the R₂ groups are independently lower alkyl, such as methyl, ethyl, propyle, butyl, and like groups, and aryl groups such as phenyl and alkylphenyl groups such as the tolyl group, as well as those where the R₂ groups are interconnected as in a divalent alkylene bridge such as



in tetrahydrothiophene oxides and dioxides. Specifically, these solvents include dimethylsulfoxide, dimethylsulfone, diphenylsulfone, diethylsulfoxide, diethylsulfone, diisopropylsulfone, tetrahydrothiophene 1,1-dioxide (commonly called tetramethylene sulfone or sulfolane) and tetrahydrothiophene-1 monoxide.

30 Additionally, nitrogen containing solvents may be used. These include dimethylacetamide, dimethylformamide and N-methyl-pyrrolidone.

The azeotrope forming solvent and polar aprotic solvent are used in a weight ratio of from about 1:10 to about 1:1, preferably from about 1:5 to about 1:3.

35 In the reaction, the hydroxy containing compound is slowly converted, *in situ*, to the alkali salt thereof by reacting with the alkali metal carbonate. The alkali metal carbonate is preferably potassium carbonate. As indicated before, mixtures of carbonates such as potassium and sodium carbonate may also be used.

Water is continuously removed from the reaction mass as an azeotrope with the azeotrope forming solvent so that substantially anhydrous conditions are maintained during the polymerization.

40 It is essential that the reaction medium be maintained substantially anhydrous during the polycondensation. While amounts of water up to about one percent can be tolerated, and are somewhat beneficial when employed with fluorinated dihalobenzoid compounds, amounts of water substantially greater than this are desirably avoided as the reaction of water with the halo and/or nitro compound leads to formation of phenolic species and only low molecular weight products are secured. Consequently, in order to secure the high polymers, the system should be substantially anhydrous, and preferably contain less than 0.5 percent by weight water during the reaction.

45 Preferably, after the desired molecular weight has been attained, the polymer is treated with an activated aromatic halide or an aliphatic halide such as methyl chloride or benzyl chloride, and the like. Such treatment of the polymer converts the terminal hydroxyl groups into ether groups which stabilize the polymer. The polymer so treated has good melt and oxidative stability.

50 While the carbonate method for preparing the polymer of this invention is simple and convenient, in some cases products of higher molecular weight can be made by the alkali metal hydroxide method. In the alkali metal hydroxide method, described by Johnson et al., U.S. Patent Nos. 4,108,837 and 4,175,175, a double alkali metal salt of a dihydric phenol is contacted with a dihalobenzoid compound in the presence of a sulfur containing solvent as herein above defined under substantially anhydrous conditions.

55 Additionally, the polymers of this invention can be prepared by other methods known in the prior art, in which at least one dihydric phenol and at least one dihalobenzoid compound are heated, for example, with a mixture of sodium carbonate or bicarbonate and a second alkali metal carbonate or bicarbonate having a higher atomic number than that of sodium, as described in U.S. Patent No. 4,176,222.

The molecular weight of the poly(aryl ethers) utilized for manufacturing the devices of the instant invention is indicated by reduced viscosity data in an appropriate solvent such as methylene chloride, chloroform, N-methylpyrrolidone, and the like. The reduced viscosities of the materials, as measured at concentrations of 0.2g per 100 ml. at 25°C, are at least 0.3 dl/g, preferably at least 0.4 dl/g and, typically, not exceeding about 5 1.5 dl/g.

The compositions of this invention are prepared by any conventional mixing method. For example, a preferred method comprises mixing the two poly(aryl ether sulfones) in powder or granular form in an extruder and extruding the mixture into strands, chopping the strands into pellets and molding the pellets into the desired article.

10 The poly(aryl ether sulfones) of the instant invention allow for the fabrication of medical devices having outstanding stress-crack resistance. These devices can be steam-sterilized under stresses of 500 psi or greater and in the presence of a variety of stream boiler additives. Typical boiler additives designated to reduce corrosion in nstream generating systems are amino compounds such as morpholine, hydrazine, N,N-diethylaminoethanol ("NALCO 359" or "BETZ NA-9"), and octadecylamine. Steam sterilization is also possible in the 15 presence of various hospital cleaners and detergents, such as those sold under the tradenames of "Castle 7900" (a sonic cleaner), "Chem Crest 14" (an ultrasonic cleaner), "Tergitol Min Foam 2X" (a non ionic surfactant), and the like.

10 The materials of the instant invention can include pigments, thermal stabilizers, ultraviolet light stabilizers, and other additives.

20 The instant poly(aryl ether sulfones) blends are useful for the fabrication for autoclave storage trays such as the systems for storage and delivery of sterile surgical instruments (thus eliminating the costs associated with wrapping); in the medical supply industry for shipment and storage of implants, prostheses and other medical devices under sterile conditions; and in many other similar applications.

25 The compositions of this invention can also be fabricated into any desired shape, i.e., moldings, coating, films, or fibers. They are particularly desirable for use as electrical insulation for electrical conductors.

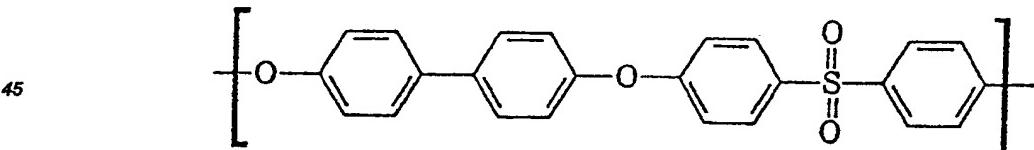
These compositions can include mineral fillers such as carbonates including chalk, calcite and dolomite; silicates including mica, talc, wollastonite, silicon dioxide, glass spheres, glass powders; aluminium; clay; quartz; and the like. Also, reinforcing fibers such as fiberglass, carbon fibers, and the like may be used. The 30 compositions may also include additives such as titanium dioxide; thermal stabilizers, ultraviolet light stabilizers, plasticizers, and the like.

EXAMPLES

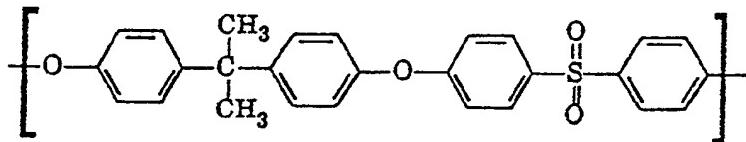
35 The following examples provide specific illustrations of the present invention and are not to be construed in any way as a limitation on its scope or generality.

Sample Preparations and Test Procedures

40 Two polymeric materials were used in all the preparation of this invention: polybiphenylsulfone and bisphenol-A polysulfone. The polybiphenylsulfone used is a polymer having the repeat unit



It is available commercially from Amoco Performance Products, Inc. under the trade name Radel R 5000. It 50 has a reduced viscosity of approximately 0.60 dl/g as measured in N-methyl pyrrolidone at a concentration of 0.2 g/dl and 25 C. It has a number-average molecular weight of about 20,000 as measured by gel permeation chromatography using methylene chloride as a solvent and a polystyrene molecular weight calibration. This polymer is referred to hereafter as PSF-I. The bisphenol-A polysulfone is the commercial product Udel P-1700, also supplied by Amoco Performance Products, Inc. It is a polymer having the following repeat unit



It has a reduced viscosity of about 0.50 g/dl in chloroform at 25°C and a concentration of 0.2 g/dl. Its number-average molecular weight by gel permeation chromatography is about 15,000 using tetrahydrofuran (THF) as solvent and a polystyrene molecular weight calibration standard. This polymer is referred to hereafter as PSF-II.

10 Both polymers were used in pellet form. Both polymers are transparent, by virtue of their amorphous character, and have a slight amber color.

Controls A and B

15 The two neat polymers PSF-I and PSF-II were dried overnight in a Lydon dehumidified recirculating air oven at a temperature of 300°F. Parts were then injection molded on a Battenfeld injection molding machine with a 3 oz injection capacity to produce standard ASTM test specimens. Mechanical properties and heat deflection temperatures (HDT) were measured per the ASTM procedures shown below. All tests were conducted on 1/8" thick specimens except HDT which was run on both 1/8" and 1/4" specimens.

20

<u>Property</u>	<u>ASTM Method No.</u>
Tensile Modulus	D-638
Tensile Strength	D-638
Elongation at Break	D-638
30 Notched Izod Impact Strength	D-256
Tensile Impact Strength	D-1822
Heat Deflection Temperature	D-648
35 Flexural Modulus	D-790
Flexural Strength	D-790

40

Examples 1 to 6

The compositions shown in Table I were mixed well as pellets and placed in a dehumidified air oven at 300°F for about 16 hrs (overnight) for drying. The dry blends were then extruded using a 25 mm twin screw double vented Berstorff extruder having an L/D ratio of 33/1 according to the conditions profile shown in Table I. The first vent port was open to the atmosphere, the second was connected to a vacuum pump. The extruder was fitted with a double strand die. The polymer extrudate was pelletized after passing through a water trough for cooling. All blends were extruded and pelletized without incident at the throughput rates indicated in Table 1. Between successive blend compositions, two pounds of extrudate were designated as "transition" material and discarded. From past experience, this amount is sufficient to effectively displace the melt in the extruder so as the compositions of the final blends do not differ from those of the dry pellet mixes.

55 The six blends were dried again overnight in the Lydon oven at 300°F and injection molded the following day on the Battenfeld injection molding machine described above to generate the needed ASTM parts. All blends appeared translucent on molding and the magnitude of the translucency increased towards the middle of the composition scale and became progressively less at the compositions rich in one polymer or the other. The lack of clarity in these blends is an indication of immiscibility between the two homopolymers.

Mechanical Properties

ASTM mechanical properties for the controls and blends are shown in Table II. The strength and modulus values for all blends are comparable to those of PSF-I. As a measure of ductility, elongation at break of all the blends can also be considered equivalent to that of PSF-I within the variability of this measurement. The tensile impact, a discriminating test of polymer unnotched toughness is likewise similar for all compositions. All these mechanical properties confirm the very good mechanical compatibility of the blends of this invention. For compositions rich in PSF-I (examples 1, 2, and 3), little or no loss in the high notched Izod of PSF-I is observed. Accordingly, compositions comprising from about 60 to 99 weight percent PSF-I and about 1 to 40 weight percent PSF-II have many of the excellent mechanical properties of the PSF-I.

Immiscible polymer blends which have poor mechanical compatibility are typically characterized by low strength for parts containing "weld lines" as a result of a multi-gated mold configuration. To ascertain the mechanical compatibility of these blends the weld-line strengths were measured against the base case with no weld line. ASTM tensile bars containing weld lines were produced on a Van Dorn injection molding machine fitted with a double gated tensile bar mold. Molding was performed at conditions similar to those used for the standard single gate case. Results are shown in Table III illustrating the retention of tensile strength over weld-line sections for all blends.

Steam Sterilization

To assess the steam sterilization resistance of the PSF-I/PSF-II blend compositions, the following test was employed. 5"x1/2"x1/8" ASTM bars were mounted by fixing the bar horizontally from one end in a cantilever fashion, and a weight was suspended on the free end. The weight attached to the specimen was such that a maximum flexural stress of about 1000 psi was generated at the fixed end surface of the bar. The samples which were mounted side by side in a stainless steel drawer were then placed in a steam autoclave. The autoclave was operated at a steam pressure of 27 psig, and a temperature of 270°F, and the autoclave chamber was charged with water containing 50 part per million (ppm) morpholine. Each autoclave cycle consisted of a 5-minute heat-up and pressurization step followed by a hold at the set conditions for 30 minutes after which the autoclave was vented and allowed to cool down for 10 minutes. The samples were typically inspected at the completion of 15-25 cycles and all samples were tested in duplicates.

Table IV summarizes the results from the steam sterilization performance study. Quite unexpectedly, the performance of all blends except that of example 6 is comparable to that of PSF-I. None showed any evidence of stress cracking following 450 cycles of steam sterilization as described above. In comparison, PSF-II experienced a rupture after only 79 cycles. The blend of example 6 had a rupture at 250 cycles - still significantly better than PSF-II. The unbroken duplicate of example 6 and control B were cracked extensively and the test was not continued on them as they are considered failures from the practical standpoint of the test. Accordingly, composition comprising from about 35 to 99 weight percent PSF-I and about 1 to 65 weight percent PSF-II have improved resistance to steam containing amine stabilizers.

Chemical Resistance

The following test was performed to evaluate the resistance of the PSF-I/PSF-II blend compositions to chemically induced stress cracking in the presence of an aqueous solution of a common nonionic wetting agent: Jet-Dry. ASTM 5"x1/2"x1/8" specimens were restrained in flexure at a constant strain on brass bend bars of two different circular curvatures to generate nominal stresses of 2,000 and 4,000 psi, respectively. The exposed surface of the molded bars was then placed in contact with the solution by partially immersing the bars into a tray containing the solution. The Jet-Dry was used at a concentration of 0.3 wt.% in water, and the test was conducted in an air oven at a temperature of 190°F for a duration of 24 hrs. The samples were inspected for physical changes following this exposure and the results are shown in Table V. This data demonstrates compositions comprising 25 to 99 weight percent PSF-I and about 1 to 75 percent PSF-II have excellent chemical resistance.

Table I: Extrusion Conditions for Preparation of PSF-I/PSF-II Blends of Examples 1-6.*

Example	1	2	3	4	5	6
Percent PSF-I	75	67	60	50	40	25
Percent PSF-II	25	33	40	50	50	75
Temperatures (C)						
Zone 1	310	310	310	280	280	280
" 2	340	340	350	315	315	315
" 3	340	340	350	330	330	330
" 4	340	340	350	340	330	330
" 5	330	340	350	340	300	300
Zone 6 and die	330	340	355	340	330	330
Melt	388	387	390	385	371	368

* Conditions common to all runs were approximately as follows:

screw speed = 200 rpm

throughput rate = 25 lbs/hr

Vent 2 vacuum reading = 30 in Hg

Vent 1 open to the atmosphere

Table II: Mechanical Properties of PSF-I, PSF-II and 6 Blend Compositions of the Two Polymers.

Example	PSF-I (%)	Tensile Strength (kpsi)	Tensile Modulus (kpsi)	Elongation at Break (%)	Flexural Strength (kpsi)	Flexural Modulus (kpsi)
Control A	100	10.3	337	109	12.8	339
1	75	10.5	344	113	13.8	363
2	67	10.4	342	100	14.1	368
3	60	10.5	357	107	14.3	370
4	50	10.4	359	99	14.5	376
5	40	10.3	364	81	14.7	379
6	25	10.3	375	96	15.1	391
Control B	0	10.0	363	132	14.8	381

5	Example	PSF-I (%)	Notched Izod (ft-lb/in)	Tensile Impact (ft-lb/in ²)	Heat Deflection Temp. [1/8"] (C)	Deflection Temp. [1/4"] (C)
10	Control A	100	16.0	235	193	203
	1	75	15.7	233	185	199
	2	67	16.1	257	184	198
15	3	60	8.1	220	179	194
	4	50	2.3	224	176	186
	5	40	1.7	211	173	186
	6	25	1.5	217	167	186
20	Control B	0	1.3	179	160	181

25 Table III: Weld Line Strengths of PSF-I/PSF-II Blends Compared with the Strength of Non-weld Line Specimens.

30	Example	PSF-I (%)	Tensile Strengths	
			No Weld Line (kpsi)	Weld Line (kpsi)
35	Control A	100	10.3	10.0
	1	75	10.5	10.3
	2	67	10.4	10.4
	3	60	10.5	10.3
40	4	50	10.4	10.4
	5	40	10.3	10.4
	6	25	10.3	10.4

45 Table IV: Steam Sterilization* Performance of PSF-I, PSF-II and 6 Blend Compositions of the Two Polymers.

50	Example	PSF-I (%)	Number of Cycles Steam Sterilization	Observations
55	Control A	100	450	No Changes
	1	75	450	No Changes

	2	67	450	No Changes
5	3	60	450	No Changes
	4	50	450	No Changes
	5	40	450	No Changes
10	6	25	250	Cracks/Rupture
	Control B	0	79	Cracks Rupture

* Steam Sterilization Conditions: 1000 psi flexural stress, 27 psig steam with 50 ppm morpholine, 275 deg F steam, 30-minute autoclaving cycle.

Table V: Chemical Resistance* of PSF-I, PSF-II and 6 Blend Compositions of the Two Polymers to Aqueous Jet-Dry (Non-ionic Based) Wetting Solution.

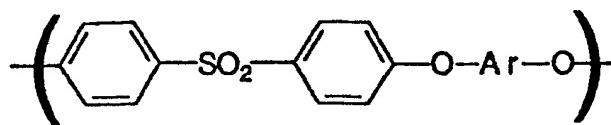
Example	PSF-I (%)	Observation at Stress Level		
		2,000 psi	4,000 psi	
Control A	100	No Changes	No Changes	
1	75	No Changes	Slight Edge Cracks	
30	2	67	No Changes	Slight Edge Cracks
	3	60	No Changes	Cracks
	4	50	No Changes	Cracks
35	5	40	No Changes	Cracks
	6	25	No Changes	Cracks
	Control B	0	Edge Cracks	Cracks

* Test Conditions: Exposure of 5"x1/2"x1/8" ASTM bars to 0.3% by weight solution at 190 deg F for 24 hrs. Flexural stress applied using constant-strain bent bars.

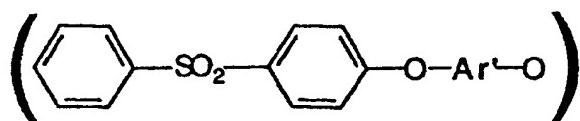
Essentially the same results can be obtained by replacing PSF-II with Ultrason S, which is the condensation of dichlorodiphenyl sulfone with approximately an 85: 15 molar ratio of Bisphenol A to Bisphenol S.

Claims

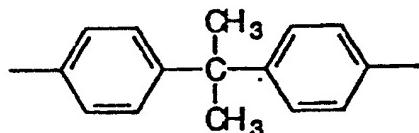
1. A composition comprising a blend of (a) poly(biphenyl ether sulfone) consisting essentially of the repeating unit:



and (b) a second poly(aryl ether sulfone) consisting essentially of the repeating unit:

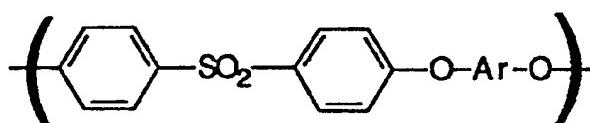


15 wherein Ar is a divalent group comprising p-biphenylene and Ar' is a divalent group comprising at least 50 mole percent

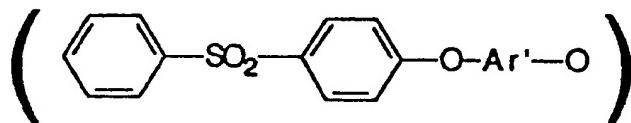


and the weight ratio of (a) to (b) is from about 25 to 99 to 75 to 1.

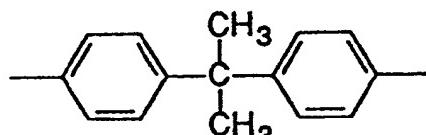
25 2. A composition comprising a blend of (a) from about 60 to about 99 percent by weight of a first poly(biphenyl ether sulfone) comprising the repeating unit



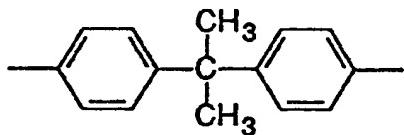
35 and (b) from about 1 to 40 percent by weight of a second poly(aryl ether sulfone) comprising the repeating unit



45 wherein Ar comprises biphenylene and Ar' comprises



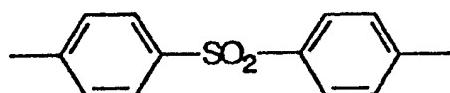
50 55 3. A composition as claimed in claim 2, wherein Ar comprises at least 50 mole percent biphenylene and Ar' comprises at least 50 mole percent



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4. A composition as claimed in claim 3, wherein from 0 to 50 mole percent of Ar and from 0 to 50 mole present of Ar' are at least one member selected from the group consisting of p-phenylene and

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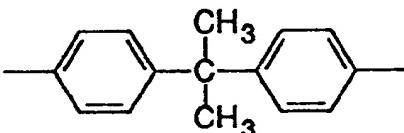


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5. A composition as claimed in claim 2, wherein Ar consists essentially of biphenylene.

6. A composition as claimed in claim 2 or claim 5, wherein Ar' consists essentially of

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7. A composition as claimed in claim 1, wherein the poly (biphenyl ether sulfone) comprises from 50 to 99 weight percent of the composition.

8. A process which comprises steam sterilizing an article formed from a composition as claimed in any one of claims 1 to 7.

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9. Use of a composition as claimed in any one of claims 1 to 7 in forming articles for medical use or for electrical insulation.

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DERWENT-ACC-NO: 1991-297566

DERWENT-WEEK: 200102

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TITLE: Poly-aryl-ether! sulphone!
compsns. for medical article
comprising poly bi:phenyl ether!
sulphone! and 2nd poly-aryl-
ether! sulphone! comprising
bisphenol=A moieties

INVENTOR: DICKINSON B L; EL-HIBRI M J ; ELHIBRI M J ; HARRIS J E ; ROBESON L M ; SAUERS M E

PATENT-ASSIGNEE: AMOCO CORP [STAD]

PRIORITY-DATA: 1990US-505826 (April 6, 1990) ,
1988US-163781 (March 3, 1988)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE
EP 450791 A	October 9, 1991	EN
CA 2038215 A	October 7, 1991	EN
JP 04227658 A	August 17, 1992	JA
US 5164466 A	November 17, 1992	EN
EP 450791 A3	July 15, 1992	EN
EP 450791 B1	May 2, 1997	EN
DE 69125882 E	June 5, 1997	DE
CA 2038215 C	July 7, 1998	EN
SG 66251 A1	July 20, 1999	EN
JP 3118011 B2	December 18, 2000	JA

DESIGNATED-STATES: DE FR GB DE FR GB

APPLICATION-DATA:

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EP 450791A	N/A	1991EP-302338	March 19, 1991
US 5164466A	N/A	1990US-505826	April 6, 1990
CA 2038215C	N/A	1991CA-2038215	March 19, 1991
DE 69125882E	N/A	1991DE-625882	March 19, 1991
EP 450791A3	N/A	1991EP-302338	March 19, 1991
EP 450791B1	N/A	1991EP-302338	March 19, 1991
JP 04227658A	N/A	1991JP-096113	April 3, 1991
JP 3118011B2	N/A	1991JP-096113	April 3, 1991
SG 66251A1	Previous Publ	1996SG-004434	March 19, 1991

INT-CL-CURRENT:

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CIPS	C08G75/23 20060101
CIPS	C08L81/00 20060101
CIPS	C08L81/06 20060101

RELATED-ACC-NO: 1989-257808 1991-297565 1994-
217092

ABSTRACTED-PUB-NO: EP 450791 A

BASIC-ABSTRACT:

Compsn. is claimed comprising: (a) poly (biphenyl ether sulphone) comprising repeating unit (I), wherein Ar is a divalent gp. comprising p-biphenylene and (b) a second poly (aryl ether sulphone) comprising repeating unit (II), wherein Ar is a divalent gp. comprising at least 50 mole % (1). The compsn. comprises 25-99 wt.% (a) and 1-75 wt.% (b). A process comprising steam sterilising an article formed from the compsn. is claimed. Use of the compsn. in forming articles for medical use or for electrical insulation is also claimed.

USE/ADVANTAGE - Poly (aryl ether sulphones) have good mechanical and thermal properties and hydrolytic stability and are widely used in medicine, their ability to be steam autoclaved is

esp. pref.. Poly (aryl ether sulphones) stress-crack when steam sterilised at 500 psi or more when excessive concns. of boiler additives e.g. morpholine are used in the steam generating system; or when contacting commonly used hospital cleansers and detergents. Poly (biphenyl ether sulphone) shows good properties but is expensive. This compsn. provides blends of poly (biphenyl ether sulphone) and a second poly (aryl ether sulphone) which, while retaining good base properties of their constituents, may be steam-sterilised at 500 psi or more, are unaffected by corrosion-reducing additives and show good chemical resistance contacting hospital cleansers and detergents. @ (13pp Dwg.No.0/0) @

TITLE-TERMS: POLY ARYL POLYETHER POLYSULPHONE
COMPOSITION MEDICAL ARTICLE COMPRISE
BI PHENYL BISPHENOL=A MOIETY

DERWENT-CLASS: A25 A85 P34 X12

CPI-CODES: A05-J06; A07-A03; A07-A03C; A11-C; A12-E01; A12-V03;

EPI-CODES: X12-E02B;

SECONDARY-ACC-NO:

CPI Secondary Accession Numbers: 1991-128650

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